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## PSYCHOSOCIAL MOOD SYMPTOMS: A COMMON PRESENTATION IN THYROID DYSFUNCTION

Rakshanda Jabeen<sup>1</sup>, Naseem Ahmed<sup>2</sup>, Muhammad Taha Tariq<sup>3</sup>, Kiran Shafiq Khan<sup>3</sup>, Bushra Perveen<sup>3</sup>, Aneeqa Khan<sup>3</sup>, Naveed<sup>3</sup>, Javaria Sohail<sup>3</sup>, Asra Waseem<sup>3</sup>, Hussaina Shabbir<sup>3</sup>, Tooba Khan<sup>3</sup>, Hafiz Abdullah<sup>3</sup>, Javeria Munir<sup>3</sup>, Bushra Ali<sup>3</sup>, Vishah Rasool<sup>3</sup>, S.M.Moiz<sup>3</sup> and A.Waqas<sup>3</sup>

<sup>1</sup>Department of Medicine, Dow Medical College, Dow University Health Sciences, Karachi, Pakistan <sup>2</sup>Departmet of Pathology, Dow Medical College, Dow University of Health Sciences, Karachi, Pakistan <sup>3</sup>Dow Medical College, Dow University of Health Sciences, Karachi, Pakistan

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## **ABSTRACT**

**Aims and objective:** This research aims at investigating the mood related consequences of thyroid dysfunction that could hint an underlying psychosocial mood disorder.

**Methods:** This cross sectional study was conducted in the Medical and Psychiatry OPD at Ruth K.M. Pfau Civil Hospital, Karachi duringOctober 2018 to May 2019. Individuals aged between 20 and 60 years, attending the outpatient department, with some nonspecific mood complain were included while those with diagnosed psychiatric or thyroid disorder were excluded. All patients were followed with the measurement of TSH, Hb, ALT, ALP and RBS.

**Results:** Among **100** participants who completed the questionnaire. Mean age of  $20 \pm 1.7$  years Mood swings and thyroid level shown a statistically significant association (p=0.047). When asked for symptoms they felt, a significant association was established between dizziness (p=0.031) and weight change (p=0.054). Additionally, thyroid levels in different age groups when compared show a remarkable association.

Conclusion: This study establishes a positive relation of thyroid dysfunction with mood disorder symptoms, hence the authors conclude that thyroid profile plays a pivotal role in the pathogenesis and management of patients with mood derangements (or an underlying psychosocial mood disorder), and is related to the overall prognosis of such patients. Much more definite work needs to be done in this domain in the future for further validation.

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## **INTRODUCTION**

Thyroid gland and its physiological role in the human metabolism plays a pivotal role, and, its disorders are some of the most researched, yet ironically have some of the most inconclusive results. Thyroid dysfunction has been directly implicated to mood derangements, and multiple psychiatric illnesses. However contradictory data exists on whether thyroid derangements are a part of nonspecific mood alterations and symptoms that could hint an underlying psychosocial mood disorder. Amongst the wide range of physiological impacts of thyroid hormones on different systems on the body, it also has been found to influence the psychiatric stability and status of an individual. Moreover, both hyperthyroidism and hypothyroidism have been found to exhibit psychiatric manifestations, including depression, mania, anxiety and, in severe derangements, psychosis as well(8, 9) Keeping in view, understandably evaluating thyroid profile in patients with psychiatric symptoms seems vital, has also proven by studies, including a study conducted in North east India by Hazarika. J et.al that elaborated the vital role of thyroid profile in management of psychiatric illness(10) Extensive research and literature is available regarding effects of thyroid on the major systems, but literature pertaining to its particular correlation with psychosocial disorders, is ambiguous. The DSM-V classified disorders, and majority have a psychosocial factor pertaining to it, including anxiety, schizophrenia, somatoform disorders and more(11). While some sources clearly disregard a correlation between thyroid and psychosocial mood disorders,(12, 13) whereas, on the contrary, others establishing a relation between thyroid dysfunction and psychosocial disorders and quality of life(11, 14)

Keeping in view this ambiguous literature, as well as paucity of literature within our local setting, the main aim of our study is to study thyroid hormone derangements in individuals presenting with symptoms of any of the psychosocial mood disorders outlined by the DSM-V, in a public hospital in Karachi.

Department of Medicine, Dow Medical College, Dow University Health Sciences, Karachi, Pakistan

<sup>\*</sup>Corresponding author: Rakshanda Jabeen

# **OBJECTIVE**

The main objective of our research is to study thyroid profile derangements in individuals presenting with symptoms of any of the psychosocial mood disorders symptoms outlined by the DSM-V, in a public hospital in Karachi.

## **METHODS**

This cross sectional study was conducted in the Medicine and Psychiatry OPD of Ruth K.M.Pfau Civil Hospital, Karachi from October 2018 to May 2019. Sample size was calculated using Open Epi software keeping anticipated frequency of 26%, gave the total sample to be **100**.

#### Inclusion Criteria

All the individuals between 20 and 60 years of age of either sex with some non-specific mood derangements, willing to partake in the study after giving viable consent.

#### Exclusion criteria

Patients already diagnosed with a psychiatric disorder, or thyroid disorders were excluded.

Moreover, patients who didn't give voluntary consent were also excluded from the study.

Participants were enrolled only after taking informed, verbal and signed consent. A face to face interview was conducted using a self-designed questionnaire. A pilot study was conducted to assess the validity of questionnaire, and correct errors in the data collection technique.

All patients were subjected to have thyroid stimulating hormone, hemoglobin, random blood sugar and alanine amino transferase and alkaline phosphatase.

Data was entered and assessed using Statistical Package for Social Sciences (SPSS) version 23. Frequencies were tabulated, and mean calculated for lab parameters. Lab parameters were compared with symptoms, using the T test, whereas categorical variables were cross tabulated using the Chi-square test. A p-value of less than 0.05 was considered significant

## **RESULTS**

Out of 100 individuals who completed the form [n=76, %] were female. Mean age among these patients was  $20 \pm 1.7$  years.

On comparing the TSH level with different symptoms with the help of Chi square, there was a significant association seen among mood swings and TSH level with a p value of 0.047. While participants feeling depressed and low showed no statically significant association (p=0.210). The rest is tabulated in Table 1.

When questioned for symptoms they felt, significant association was found between dizziness (p=0.031) and weight change (p=0.054) (Table 1).

We further extend our result by comparing thyroid levels in different age groups, shows a significant relation when compare as in table  $2\,$ 

To further describe our result we compare TSH level with all other labs test results, where no significant association was recognized but an inverse relation was built between TSH and hemoglobin (p=0.176) level also with Alkaline phosphatase (p=0.325) showing an insignificant relation too. (Table 3)

**Table 1** Responses of participates of different thyroid level with different symptoms

		,	1		
		Hypothyroi dism	Normal	Hyperthyroi dism	p-value
Has there been any major	Yes	0	5 (100%)	0	0.476
life events in previous 3					
months (birth, death, divorce)	No	4 (4.2%)	73 (76.5%)	18 (18.9%)	
Have you had any mood	Yes	2 (2.6%)	65 (83.3%)	11 (14.1%)	0.047
swings recently	No	2 (9.1%)	13 (59.1%)	7 (31.8%)	
Do you feel depressed or	Yes	2 (2.6%)	60 (76.8%)	16 (20.5%)	0.210
	hope	less on trivial r	natters		
	No	2 (9.1%)	18 (8	1.8%)	2 (9.1%)
D	Yes	3 (3.3%)	70 (77.8%)	17 (18.9%)	0.496
Do you often feel anxious	No	1 (10%)	8 (80%)		1 (10%)
Do you feel tired or	Yes	3 (3.4%)	66 (75.9%)	18 (20%)	0.166
lethargic all the times	No	1 (7.7%)	12 (9.3%)		0
Do you feel difficulty in	Yes	1 (1.6%)	48 (78.7%)	12 (19.7%)	0.298
enjoying things do you often lose your temper	No	3 (7.7%)	30 (76.9%)		6 (15.4%)
Do you have any difficulty	Yes	0	43 (86%)	7 (14%)	0.058
in recalling things	No	4 (8%)	35 (70%)	11 (22%)	
Have you ever had any	Yes	0	13 (86.7%)	2 (13.3%)	0.580
suicidal/ homicidal thoughts	No	4 (4.7%)	65 (76.5%)	16 (18.8%)	
Do you have any difficulty	Yes	2 (6.5%)	24 (77.4%)	5 (16.1%)	0.685 13
in concentration	No	2 (2.9%)	54 (78.3%)		(18.8%)
1		feel any symp			
Fainting	Yes	1 (3.4%)	24 (82.8%)	4 (13.4%)	0.758
1 umung	No	3 (4.2%)	54 (76.1%)	14 (19.7%)	
Palpitation	Yes	1 (1.6%)	49 (80.3%)	11 (18.0%)	0.319
1 dipitation	No	3 (7.7%)	29 (74.4%)	7 (17.9%)	
Generalized body ache	Yes	3 (3.3%)	72 (78.6%)	17 (18.5%)	0.422
Generalized body acid	No	1 (12.5%)	6 (75%)	1 (12.5%)	
Illness induce anxiety	Yes	0	17 (77.7%)	5 (22.7%)	0.477
muce unitely	No	4 (5.1%)	61 (78.9%)	13 (16.7%)	
Dizziness/ light headache	Yes	1 (1.3%)	63 (81.8%)	13 (16.9%)	0.031
Dizziness/ fight fleatache	No	3 (13.0%)	15 (65.2%)	5 (21.7%)	
Alter sleep	Yes	1 (1.9%)	42 (77.8%)	11 (20.8%)	0.423
Atter steep	No	3 (6.5%)	36 (78.5%)	7 (15.2%)	
Alter bowel habit	Yes	2 (6.7%)	23 (76.6%)	5 (16.5%)	0.666
Auto bowei nabit	No	2 (2.9%)	55 (78.9%)	13 (18.3%)	
Weight change	Yes	2 (7.4%)	24 (88.9%)	1 (3.7%)	0.054
weight change	No	2 (2.7%)	54 (74.0%)	17 (23.3%)	

**Table 2** comparing variable thyroid level with different age groups

Age	Hypothyroidism	Normal	Hyperthyroidism	p-value
20-30	0	4 (87.5%)	2 (12.5%)	0.039
31-45	0	0	35 (83.3%)	7 (16.7%)
46-60	2 (5.9%)	2 (5.9%)	24 (70.9%)	8 (23.5%)
>60	2 (25%)	2 (25%)	5 (62.5%)	1 (12.5%)

**Table 3** Correlation of TSH level with other Lab parameters

Independent variable	Dependent variable	Co-relation	p-value
TSH level	RBS level	0.192	0.149
TSH level	Hemoglobin level	-0.142	0.176
TSH level	Alkaline phosphatase	-0.197	0.325
TSH level	ALT level	0.070	0.552

## **DISCUSSION**

The clinical implications of thyroid hormones in mood disorder have been studied extensively and still remains disputable (15). Even though thyroid disorders are associated with psychiatric symptoms in clinical populations, extent of a similar association in general population is less certain .A group of investigators at Copenhagen conducted prospective cohort studies to determine the association between thyroid and affective disorders, giving positive results (16, 17). in a research conducted in India there was an intimate relation between Mood disorders and suboptimal thyroid function too Moreover. the most recent ultrasensitive immunoradiometric assay for detecting thyroid dysfunction is TSH (19). Therefore, Our clinical trial also uses TSH levels for establishing a relation with mood swings.

Overt or subclinical Hypothyroidism, appears to the commonest abnormality found in bipolar disorder. The most common psychiatric indication related to hypothyroidism are depression and cognitive dysfunction (18). Both the symptoms of hypothyroidism and hyperthyroidism may be presented with various neuropsychiatric manifestations including mild depression disorders, anxiety and overt psychosis. In our study it is clearly shows that apart from majority of euthyriod patients, more patients presented with hyperthyroidism as compared to the number of hypothyroid patients. There was no statistically significant relation between major life event such as divorce, death or birth. While mood swings and thyroid level has shown a statistically significant association. Difficulty in recalling things, dizziness/light headedness and weight changes also shows significant results. This shows that disturbance in TSH levels can lead to affective disorders such as bipolar disorder

Male patients presented significantly higher TSH levels and better clinical improvement (22). In contrast to their study, our randomized clinical trial surprisingly shows that higher TSH levels were found in females patients (18.4%) as compared to males (16.7%). Whereas low TSH levels (hypothyroidism) were more prevalent in males (4.2%) as compare to females (3.9%).So the authors hypothesize that gender related differences of TSH levels may be connected with different metabolism and requires further investigation. (Refer to table 1). In our population levels of TSH was between references values in each individual, so far undiagnosed, thyroid disorder were not apparent.

Biondi B *et al.* reported that the commonness of thyroid disorders rises with age. They become common in individuals aged 60 years and older (23). In contrast to their findings, our study shows that all the age groups presented with hyperthyroidism more commonly with the exception of age group 60 and above , where hypothyroidism is more prevalent(25%). Among the hyperthyroid patients, the most pervasive age group was 46-60 years (23.5%) while in the same age group the hypothyroid patients were just 5.9%. (Refer to table 5).

A study by Hong w j *et al.* shows that a pattern of euthyriod hyperthyroxinemia is significantly more common in patients with a mood disorder and in Colorado, thyroid disease prevalence study shows that low mood disorder is a poor predictor of hypothyroidism (26) which is consistent with the results of our study in which there was a statistically significant relation between mood swings and thyroid symptoms. We further investigated that the symptom of mood swings were more prevalent in the patient of hyperthyroidism (14.1%) as compared to hypothyroidism (2.6%) (27).

A study conducted by d fang *et al.* found patients with hyperthyroidism had executive function and could not inhibit impulsive behavior (28), but Vogels study revealed no cognitive impairment in patients with Graves's thyrotoxicosis using comprehensive neuropsychological testing. Emotional disturbance of patients with acute untreated hyperthyroidism have been reported in early time (29). Our results were contrary to these studies, where patients of hyperthyroidism had more affective symptoms both of depression (20.5%) and anxiety (18.9%).

In a study at UK, aimed at evaluating the mental health in general practice, patients referred for thyroid function test had

a higher rate of psychological morbidity than seen in all presentations (54.2 vs. 19%), suggesting that psychiatric symptoms were a common reason for assessing thyroid function (30) Contrary to this studies, our results clearly show that there is no statistically consequential relation

#### Limitations and Future Recommendations

This research according to the best of our knowledge is the first of its kind in our setup and has attempted to explain the complex relationships among the symptoms of mood disorders and TSH levels in addition to the relationship between TSH and levels of ALT, RBS and hemoglobin, The main strength lies in the fact that authors did not solely rely on the patients symptomatology, but included lab investigations for a broader explanation of the results.

These include the usage of self-administered questionnaire instead of a standard one. An important limitation in this regard is differences in thyroid hormone metabolism between male and female patients and low sample size may have confounded our results. Although current research, especially preclinical, research has provided strong leads it is further recommended that more factors should be incorporated to find out the probable relation between the two. The authors also recommend that future researches should be done with a higher sample size and should necessarily incorporate both free t4 and t3 levels in addition to TSH levels for better understanding of the possible complex interplay between these hormones and mood disorders. The authors also recommend that future researches in this regard should have an equal number of male and female participants, which will lead to more authentic results about the relationship between gender and thyroid problems.

# **CONCLUSION**

This study reveals a statistically significant relation between mood swings and thyroid levels. The researchers of this study conclude that mechanisms, by which thyroid dysfunction produces mood symptoms, remain to be more fully elaborated and understood. Studies among depression are more prevalent than those of mood disorder in our set up and that more research should be done in this subject. Therefore, more methodologically stable studies among clinical studies are required to assess potential interactions between these neurochemical systems in the CNS and thyroid functions.

## References

- 1. Bauer M, Goetz T, Glenn T, Whybrow PC. The thyroid-brain interaction in thyroid disorders and mood disorders. J Neuroendocrinol. 2008;20(10):1101-14.
- 2. Breitzig MT, Alleyn MD, Lockey RF, Kolliputi N. Thyroid hormone: a resurgent treatment for an emergent concern. Am J Physiol Lung Cell Mol Physiol. 2018;315(6):L945-L50.
- 3. Szlejf C, Suemoto CK, Santos IS, Lotufo PA, Haueisen Sander Diniz MF, Barreto SM, *et al.* Thyrotropin level and cognitive performance: Baseline results from the ELSA-Brasil Study. Psychoneuroendocrinology. 2018;87:152-8.
- 4. Hall JE. Guyton and Hall textbook of medical physiology. 11 ed. Philadelphia, PA: Elsevier; 2016.
- 5. Moriggi G, Verga Falzacappa C, Mangialardo C, Michienzi S, Stigliano A, Brunetti E, *et al.* Thyroid

- hormones (T3 and T4): dual effect on human cancer cell proliferation. Anticancer Res. 2011;31(1):89-96.
- Moeller LC, Fuhrer D. Thyroid hormone, thyroid hormone receptors, and cancer: a clinical perspective. Endocr Relat Cancer. 2013;20(2):R19-29.
- 7. Hage MP, Azar ST. The Link between Thyroid Function and Depression. Journal of thyroid research. 2012;2012:590648-.
- 8. Hage MP, Azar ST. The Link between Thyroid Function and Depression. J Thyroid Res. 2012;2012:590648.
- Bauer M, Berman S, Stamm T, Plotkin M, Adli M, Pilhatsch M, et al. Levothyroxine effects on depressive symptoms and limbic glucose metabolism in bipolar disorder: a randomized, placebo-controlled positron emission tomography study. Mol Psychiatry. 2016;21(2):229-36.
- Hazarika J, Kalita KN, Sharma M, Saikia S, Patangia P, Hazarika P, et al. Thyroid profile in depression: a crosssectional study from North-East India. 2017;5(3):1066.
- Psychosocial Personality Disorders symptoms, meaning, average, Definition, Description, Demographics, Causes and symptoms [Available from:http://www.healthofchildren.com/P/Psychosocial-Personality-Disorders.html.
- 12. Kritz-Silverstein D, Schultz ST, Palinska LA, Wingard DL, Barrett-Connor E. The association of thyroid stimulating hormone levels with cognitive function and depressed mood: the Rancho Bernardo study. J Nutr Health Aging. 2009;13(4):317-21.
- 13. Menon B. Hypothyroidism and bipolar affective disorder: is there a connection? Indian J Psychol Med. 2014;36(2):125-8.
- 14. Teixeira PdFdS, Reuters VS, Almeida CP, Ferreira MM, Wagman MB, Reis FAA, *et al.* Evaluation of clinical and psychiatric symptoms in sub clinical hypothyroidism. 2006;52(4):222-8.
- 15. Gibney SM, Drexhage HAJJoNP. Evidence for a dysregulated immune system in the etiology of psychiatric disorders. 2013;8(4):900-20.
- 16. Thomsen AF, Kessing LVJBd. Increased risk of hyperthyroidism among patients hospitalized with bipolar disorder. 2005;7(4):351-7.

- 17. Thomsen AF, Kvist TK, Andersen PK, Kessing LVJEJoE. Increased risk of affective disorder following hospitalisation with hyperthyroidism–a register-based study. 2005;152(4):535-43.
- 18. Larsen J, Faber J, Christensen E, Bendsen B, Solstad K, Gjerris A, *et al.* Relationship between mood and TSH response to TRH stimulation in bipolar affective disorder. 2004;29(7):917-24.
- 19. Bschor T, Baethge C, Adli M, Lewitzka U, Eichmann U, Bauer MJJoP, *et al.* Hypothalamic-pituitary-thyroid system activity during lithium augmentation therapy in patients with unipolar major depression. 2003;28(3):210.
- 20. Taylor JWJTAjop. Depression in thyrotoxicosis. 1975.
- 21. Whybrow P, Prange A, Treadway CJAoGP. Mental changes accompanying thyroid gland dysfunction: a reappraisal using objective psychological measurement. 1969;20(1):48-63.
- Aronson R, Offman HJ, Joffe RT, Naylor CDJAogp. Triiodothyronine augmentation in the treatment of refractory depression: a meta-analysis. 1996;53(9):842-8
- 23. Biondi B, Cooper DSJEr. The clinical significance of subclinical thyroid dysfunction. 2007;29(1):76-131.
- 24. Joffe RT, Pearce EN, Hennessey JV, Ryan JJ, Stern RAJIjogp. Subclinical hypothyroidism, mood, and cognition in older adults: a review. 2013;28(2):111-8.
- 25. Shashi A, Sharma NJIJoB, Sciences AM. Prevalence and clinical aspects of thyroid disorders in himachal pradesh, india. 2015;5(1):86-94.
- Canaris GJ, Manowitz NR, Mayor G, Ridgway ECJAoim. The Colorado thyroid disease prevalence study. 2000;160(4):526-34.
- 27. Dickerman AL, Barnhill JWJAJoP. Abnormal thyroid function tests in psychiatric patients: a red herring? 2012;169(2):127-33.
- 28. Dai F, Yuan L, Fang J, Zhang Q, Wang KJNl. Impaired decision making under risky conditions in the acute phase of Graves' thyroitoxicosis. 2017;661:1-4.
- 29. Vogel A, Elberling TV, Hørding M, Dock J, Rasmussen ÅK, Feldt-Rasmussen U, *et al.* Affective symptoms and cognitive functions in the acute phase of Graves' thyrotoxicosis. 2007;32(1):36-43.
- 30. Dayan CM, Panicker VJEtj. Hypothyroidism and depression. 2013;2(3):168-79.

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